IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Docket No.: 014835-101.02-029 In re United States Patent Application of: Conf. No.: 6557 **Applicants:** Geddes, et al. 10/536,502 Art Unit: 2858 Application No.: Examiner Angela M Bertagna Date Filed: 12/14/2005 Title: HIGH SENSITIVITY ASSAYS Customer No.: 24239 FOR PATHOGEN **DETECTION USING METAL-**ENHANCED FLUORESCENCE :

DECLARATION OF INVENTORS IN U. S. PATENT APPLICATION NO. 10/536,502

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

Sir:

CHRIS GEDDES, JOSEPH R. LAKOWICZ and LES BAILLIE severally and jointly hereby declare:

- THAT We are co-inventors of the subject matter disclosed and claimed in United States Patent Application No. 10/536,502, filed December 14, 2005 in the United States Patent and Trademark Office entitled, "HIGH SENSITIVITY ASSAYS FOR PATHOGEN DETECTION USING METAL-ENHANCED FLUORESCENCE," claiming priority from PCT International Application No.: PCT/US03/38163 filed on November 26, 2003 which in turn claims priority to US Provisional Application No 60/429,263 filed on November 26, 2002, hereafter referred to as the "Application."
- 2) THAT the Application discloses and claims a system that includes the use of two separate and distinct probe sequences (a captured and free nucleotide sequence probes) that are not complementary to each other but instead are complementary to separate and distinct portions of

the target nucleotide sequence, which is anthrax. Importantly, with the specific use of two probes sequences that are complementary to different sections of the target nucleotide sequence, the free probe sequence includes a fluorophore that is positioned a distance from the metallic surface to provide for a more sensitive test with increased reliability even when the level of anthrax in the sample is not identifiable by other less sensitive tests.

- 3) THAT We are aware that the Application has been examined by the United States Patent and Trademark Office and aware that the claims of the Application have been rejected on various grounds including the disclosure of Cao, et al., Nanoparticles within Raman Spectroscopic Fingerprints for DNA and RNA Detection, Science, Aug 2002, Vol. 297, pp 1536-1540, hereinafter Cao.
- 4) THAT we have been informed by my legal representative that the rejections of the claims of the Application can be overcome by presenting evidence to the United States Patent and Trademark Office of our possession of the claimed invention prior to the August 30, 2002 publication date of the Cao reference (Publication Date).
- 5) THAT attached is Exhibit 1 hereof which is a true and exact copy of power point slide, with dates blackened out, that was created prior to the Publication Date for inclusion in a grant proposal. The slide clearly shows the use of two separate nucleotide sequence probes that have affinity for different sections of a target sequence of anthrax. Further, this slide shows that one of the probes may contain not only a fluorophore but also include a metal particle. Further included is Exhibit 2 which is a copy of the Disclosure and Record of Invention that was submitted after the Publication Date of Cao, where in the document states that the invention was reduced to practice. The provisional application relating to this invention was timely filed by November 26, 2002.
- 6) THAT we offer Exhibits 1 and 2 with this Declaration as evidence of the conception, diligence by us the inventors of the anthrax assay system of the present invention prior to the Publication date of August 30, 2002 and reduction to practice after the Publication date with a subsequent filing of a provisional application on November 26, 2002.

As a below-named declarant, I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements, and the like, so made are punishable by fine or imprisonment, or both, under Section 1001 or Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Chris Geddes PhD
Date
Ind ner
Joseph R. Kakowicz, PhD
april 29, 2009
Date
Les Baillie, PhD
Date

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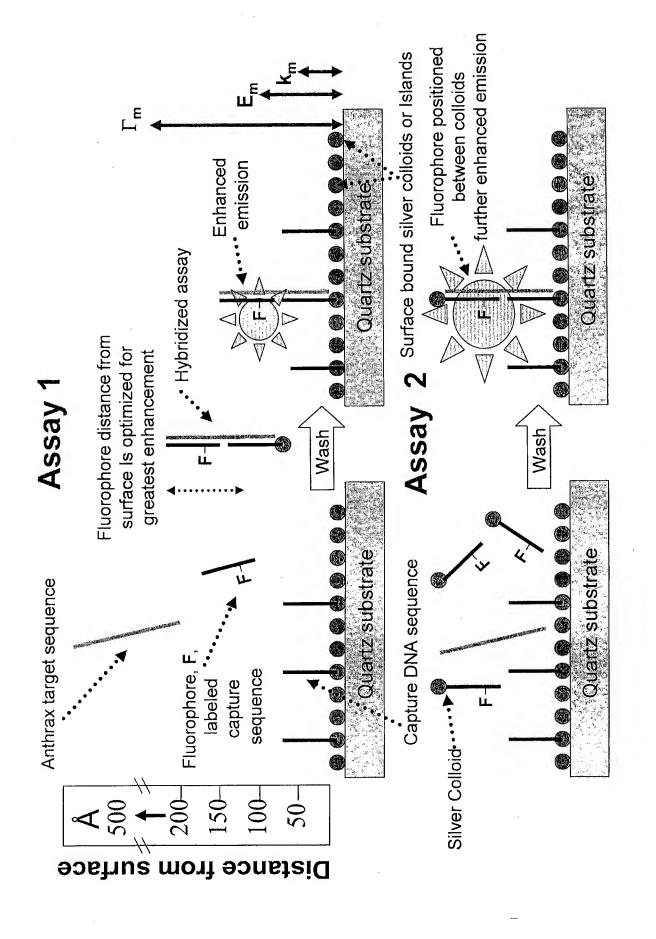
As a below-named declarant, I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements, and the like, so made are punishable by fine or imprisonment, or both, under Section 1001 or Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Chris Geddes PhD	
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Date	
Joseph R. Lakowicz, PhD)
Date	
L Rux	
Les Baillie, PhD	0
16th APRIL 2006	
Date	

As a below-named declarant, I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements are made with the knowledge that willful false statements, and the like, so made are punishable by fine or imprisonment, or both, under Section 1001 or Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

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Chris Geddes PhD	
4-28-09 Date	
Joseph R. Lakowicz, PhD	and a second
Date	
Les Baillie, PhD	· · · · · · · · · · · · · · · · · · ·
Date	

Exhibit 1



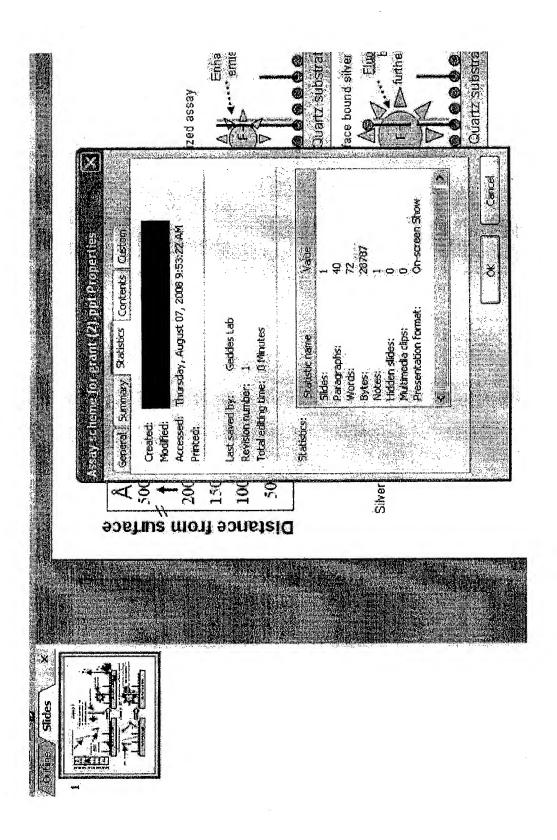


Exhibit 2

UNIVERSITY OF MARYLAND BIOTECHNOLOGY INSTITUTE (UMBI)

UMBI File No. 02-029-J-DC

Date Received

DISCLOSURE AND RECORD OF INVENTION

This invention was made and/or reduced to practice in the course of work performed at the UMBI facility.

Mail completed disclosure to UMBI Patent Department, 701 E. Pratt Street, Suite 200, Baltimore, MD 21202

Telephone (410) 385-6324-25, if you have questions regarding this form.

Telephone (410) 385-6324-25, if you have questions regarding this form.					
<u>Title of Invention:</u> High-Sensi	tivity Assays for Path	nogen Detection using	g Metal-Enhan	ced Fluorescend	ce .
Please list names of those who contributiled the Patent Department will make	ated to the inventive the legal determination o	ought and to the final refinventorship.	sult of the inver	ntion. If a patent a	pplication is
Contributor(s): Full Name	Title/Position	Employer	<u>Division</u>	Phone No.	Mail Stop
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(40 %) Joseph R. Lakowicz	Professor	UMAB		ne (410) 706 4809	<u>.</u>
(20 %) Les Baillie	Assoc. Professor	UMBI	MBC	(410) 706 4565	5
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Chris D. Geddes	UK	15 Charles Plaza,	#805,	Baltimore	MD, 21201
Les Baillie	UK	7356 Eden Brook Drive, #715,		Columbia, MD, 21046.	
Joseph R. Lakowicz	USA	10037 Fox Den Roa	<u>d</u>	Ellicott City, 1	MD, 21042
Government Grants Funding Agency and Grant No			is invention		
Principal Investigator Sponsored Research Grants Sponsor Name Principal Investigator					
Private, Non-Government Grants Funding Agency and Grant No. Principal Investigator Subcontract Subcontract No. Contractor					
Other Describe				UMBI	
From what source do you reasonably expect future or continuing funding?					
Source NIH		Is this c	ontinuing fund	ding or X future	funding?
No future funding is currently	expected. We ha	ve just submitted an l	R01 proposal	to the NIH (attac	ched).

Conception (Date, Place):	
Earliest documentation of	your invention: (please provide date and identify the document)
First Sketch or Drawing:	Not Disclosed
First Written Description	: Not Disclosed
Names of witnesses or o	others with knowledge of facts relating to conception:
Inventors	
Reduction to Practice:	
Date first model complete	ed:
Date of operation and te	sting: See 1 on attached sheet
Place of test:	
Results of testing:	
Witnesses or others with	n direct knowledge of test:
Invantava	
Inventors	
Important: WAS ANY PROPRII (e.g., software code	ETARY MATERIAL FROM OUTSIDE YOUR LABORATORY USED TO DEVELOP YOUR INVENTION a, cell line, antibody, DNA fragments, or chemical compound)? If yes, please describe.
(2.0.) 2.3	
No	
DESCRIPTION:	
What your invention er	Please summarize: ercome to make the invention, nables people to do that couldn't be done as well before, address the problem your invention addresses.
	ń.
	y ir is
	See 2
Summary of the Invention	Please include a sketch of the invention if possible (you may attach extra pages):
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	See 3
	Dec 3

				
List uses of the invention - research, commercial, pilot plant, etc.; Think as broadly as possible. Identify companies that might be interested in licensing this technology.				
	See 4			
Publications (including UMBI reports, m DESCRIBE or RELATE to the invention:	eeting abstra	cts, or prior patents or p	patent applications) that <u>ACTUALLY</u>	
<u>Title/Subject</u>	Publication Date	Indicate "Describes" or "Relates" to Inv.	Journal and/or UMBI Rep. No. If prior UMBI pat. appl., Disclosure ID#	
See attached list of patents and papers		Relates	-	
which relate to this invention disclosure	- -			
Papers you have SUBMITTED or are PRI	<u>EPARING</u> for p	ublication that describe	or relate to your invention:	
<u>Title/Subject</u>	Anticipated Publ. Date	Indicate "Describes" or "Relates" to Inv.	Journal and/or UMBI Rep. No. If prior UMBI pat. appl., Disclosure ID #	
No papers submitted or pending. Howe	ver we have att	ached a recently submitte	d R01 grant proposal based on this disclosure.	
Please notify the Patent Dept. (x 6324-2	5) if you later	plan new publications	that describe or relate to the invention.	

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I believe myself to have contributed to be the above-described invel	ntion. (Each contributor must sign.)
CONTRIBUTOR: (40% Contribution)	DATE:
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WITNESS: Man losufall	DDATE:
CONTRIBUTOR: (20% Egntribution) Pr I balle	DATE:
WITNESS: Man localella	DATE:
Read and Understood by:	
UMBI Patent Department Reviewer	Date:
Center's Director	Date: